

INDIVIDUALIZED INTRAFIBER CROSSLINKED CELLULOSIC FIBERS WITH IMPROVED BRIGHTNESS AND COLOR

FIELD

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The present application relates to individualized intrafiber crosslinked cellulosic fibers with improved color and brightness properties.

BACKGROUND

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Cellulosic fibers are a basic component of absorbent products such as diapers. The ability of an absorbent product containing cellulosic fibers to initially acquire and distribute liquid will generally depend on the product's dry bulk and capillary structure. However, the ability of a product to acquire additional liquid on subsequent insults will depend on the product's wet bulk. Cellulosic fibers, although absorbent, tend to collapse on wetting and to retain absorbed liquid near the point of liquid insult. The inability of wetted cellulosic fibers in absorbent products to further acquire and distribute liquid to sites remote from liquid insult can be attributed to a diminished acquisition rate due in part to the loss of fiber bulk associated with liquid absorption. Absorbent products made from cellulosic fluff pulp, a form of cellulosic fibers having an extremely high void volume, lose bulk on liquid acquisition and the ability to further wick and acquire liquid, causing local saturation.

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Crosslinked cellulosic fibers generally have enhanced wet bulk compared to uncrosslinked fibers. The enhanced bulk is a consequence of the stiffness, twist, and curl imparted to the fiber as a result of crosslinking. Accordingly, crosslinked fibers are advantageously incorporated into absorbent products to enhance their wet bulk and liquid acquisition rate and to also reduce rewet.

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Some of the first crosslinked cellulosic fibers were prepared by treating cellulosic fibers with formaldehyde and various formaldehyde addition products. See, for example, U.S. Patent No. 3,224,926; U.S. Patent No. 3,241,553; U.S. Patent No. 3,932,209; U.S. Patent No. 4,035,147; and U.S. Patent No. 3,756,913. Unfortunately, the irritating effect of formaldehyde vapor on the eyes and skin is a marked disadvantage of the fibers. In addition, such crosslinked fibers typically exhibit objectionable odor and have low fiber brightness.

Alternatives to formaldehyde and formaldehyde addition product crosslinking agents have been developed. Among these are dialdehyde crosslinking agents. See, for example, U.S. Patent No. 4,822,453, which describes absorbent structures containing individualized, crosslinked fibers, wherein the crosslinking agent is selected from the group consisting of C₂-C₉ dialdehydes, with glutaraldehyde being preferred. The reference appears to overcome many of the disadvantages associated with formaldehyde and/or formaldehyde addition products. However, the cost associated with producing fibers crosslinked with dialdehyde crosslinking agents such as glutaraldehyde is considered too high to result in significant commercial success. Therefore, further efforts have been made to improve fiber properties such as color and odor.

Polycarboxylic acids have been used to crosslink cellulosic fibers. See, for example, U.S. Patent No. 5,137,537; U.S. Patent No. 5,183,707; and U.S. Patent No. 5,190,563. These references describe absorbent structures containing individualized cellulosic fibers crosslinked with a C₂-C₉ polycarboxylic acid. The ester crosslink bonds formed by the polycarboxylic acid crosslinking agents differ from the acetal crosslink bonds that result from the mono- and di-aldehyde crosslinking agents. Absorbent structures made from these individualized, ester-crosslinked fibers exhibit increased dry and wet resilience and have improved responsiveness to wetting relative to structures containing uncrosslinked fibers. Furthermore, the preferred polycarboxylic crosslinking agent, citric acid, is available in large quantities at relatively low prices making it commercially competitive with formaldehyde and formaldehyde addition products. Unfortunately, the preferred C₂-C₉ crosslinking agent, citric acid, can cause discoloration (i.e., yellowing) of the white cellulosic fibers when the treated fibers are cured at the elevated temperatures required for crosslinking. It is known that decomposition of citric acid yields aconitic acid, itaconic acid, citraconic, and mesaconic acid. Yellowing may be due to the chromophores

produced as a result of the conjugated double bonds produced or due to reactions with the double bonds. In addition, unpleasant odors can also be associated with the use of α -hydroxy polycarboxylic acids such as citric acid. The above-noted references do not describe processes that reduce the odor or increase the brightness of the treated fibers.

5 More recently, it was found that the characteristic odor associated with citric acid crosslinked cellulosic fibers could be reduced and the brightness improved by contacting the fibers with an alkaline solution (e.g., an aqueous solution of sodium hydroxide) and an oxidizing bleaching agent (e.g., hydrogen peroxide). See U.S. Patent No. 5,562,740. In the method, the alkaline solution raises the finished fiber pH preferably to the
10 5.5-6.5 range from about 4.5. This in combination with the oxidizing bleaching agent reduces the "smoky and burnt" odor characteristics of the crosslinked fibers. The oxidizing bleaching agent when added at high consistency increases the final product brightness to 80 to 86 from 70 to 75 and reduces odor.

Although some disadvantages related to brightness and color associated with crosslinked
15 cellulosic fibers have been addressed, a need remains for cellulosic fibers having the advantages of bulk, liquid acquisition, and rewet associated with crosslinked cellulosic fibers without the disadvantages related to diminished fiber brightness and color. The present application seeks to fulfill these needs and provides further related advantages.

SUMMARY

20 In one aspect, individualized cellulosic fibers having improved color and brightness are disclosed. The cellulosic fibers are intrafiber crosslinked cellulosic fibers obtainable from cellulosic fibers by treatment with a crosslinking agent in the presence of a polyol. The fibers have a brightness greater than about 79.0 % ISO, color characterized by a Whiteness Index

25 $(WI_{(CDM-L)})$ greater than about 69.0 and bulk greater than about 16 cm³/g.

In another aspect, methods for the preparation of cellulosic fibers having improved brightness and color are provided. In the methods, a fibrous web of cellulosic fibers is treated with a crosslinking agent in the presence of a polyol and then cured to provide individualized cellulosic fibers having improved brightness and color.

30 In still another aspect, absorbent products are provided incorporating the crosslinked fibers with improved color and brightness.

DETAILED DESCRIPTION OF THE EMBODIMENTS

The present application provides cellulosic fibers having improved brightness and color and methods for their preparation. The fibers are intra-fiber crosslinked cellulosic fibers which have been treated with a crosslinking agent in the presence of a polyol. The crosslinked cellulosic fibers are made under pilot plant conditions representative of commercial production by treatment with an effective amount of a crosslinking agent and an amount of polyol effective to provide crosslinked fibers having a brightness greater than about 79.0 % ISO. The polyol treated fibers which have been crosslinked have a color characterized by an Whiteness Index ($WI_{(CDM-L)}$) value greater than about 69.0, an L value greater than about 94.5, an a value greater than about -1.55 and less than about -0.60, and a b value less than about 8.5.

The term “polyol” means “a polyhydric alcohol, i.e., one containing three or more hydroxyl groups”. Those having three hydroxyl groups (trihydric) are glycerols; those with more than three are called sugar alcohols, with general formula $CH_2OH(CHOH)_nCH_2OH$, where n may be from 2 to 10. Pigman in “The Carbohydrates”, W. Pigman, Editor, Academic Press Inc., NY, 1957, divides polyols into two classes, the acyclic polyols (alditols, glycitols, or “sugar alcohols”) and the alicyclic polyols (cyclitols). The term “polyol” as used in this application also includes heterosides which contain a single polyol linked by a glycosidic bond to another carbohydrate. Additionally, the polyol may be linked to one, two, or three sugars. Examples include, but are not limited to lactitol, mannitol and isomalt. The acyclic polyols include, but are not limited to, triitol which includes glycerol; tetrیتols including threitol and erythritol; pentitols including arabinitol, xylitol, ribitol, rhamnitol and fucitol; hexitols include sorbitol, mannitol, talitol, iditol, galactitol, and allitol; heptitols include volemitol, perseitol, β -sedoheptitol, *D-glycero-D-ido*-heptitol, *meso-D-glycero-L-ido*-heptitol and siphulitol; octitols include *D-erythro-L-gala*-octitol, *D-erythro-D-gala*-octitol, *erythro-manno*-octitol, *D-erythro-L-talo*-octitol, *D-threo-L-gala*-octitol; α,α,α -*D-Gluco*-nonitol; $\alpha,\alpha,\alpha,\alpha$ -*D-Gluco*-decitol. The alicyclic polyols are polyhydroxy derivatives of cyclohexane and include *cis*-Inositol, *epi*-Inositol, *allo*-Inositol, *neo*-Inositol, *myo*-Inositol, 1*D-chiro*-Inositol, 1*L-chiro*-Inositol, *muco*-Inositol, and *scyllo*-Innositol. Alicyclic polyols with only four hydroxyl groups include betitol, *L-leucanthemitol* and conduritol; cyclic polyols with five hydroxy groups include *D-*

quercitol, L-quercitol and L-viburnitol. The heterosides include lactitol, maltitol, and isomalt. The latter consists of two components in a 1:1 mixture, 6-O- α -D-Glucopyranosyl-D-sorbitol and 1-O- α -D-Glucopyranosyl-D-mannitol. Others include clusianose, umbilicin and peltigeroside.

5 The crosslinked cellulosic fibers with improved brightness and color properties are made by treating a mat or web of cellulosic fibers with an aqueous solution of a crosslinking agent in the presence of the polyol to provide treated fibers, which are then separated into individually treated fibers, and heated for a time and at a temperature to effect drying and subsequently curing (i.e., to provide intrafiber crosslinked cellulosic fibers with improved
10 brightness and color). A representative method for making the crosslinked cellulosic fibers with improved color and brightness properties is described in Example 1. The term "brightness" refers to the reflectance of blue light corresponding to a centroid wavelength of 457 nm in terms of the perfect reflecting diffuser (perfect reflecting diffuser is the ideal reflecting surface that neither absorbs nor transmits light, but reflects
15 diffusely, with the radiance of the reflecting surface being the same for all reflecting angles, regardless of the angular distribution of the incident light). Brightness was measured according to TAPPI T 525 om-02 on a Technibrite MicroTB-1C instrument (Technydine Corp.).

In one embodiment, the crosslinked fibers have a brightness greater than about 79.0 %
20 ISO. The brightness and color properties of fibers as a function of the type of polyol in the presence of a crosslinking agent are presented in Tables 1 and 2. Table 3 represents the effect of time and temperature under pilot plant conditions which are representative of commercial production.

In addition to high brightness, the crosslinked fibers prepared with the polyols exhibit
25 improved color properties as indicated by the Opponent colors scales L , a , b values, (Hunter space), and Whiteness Index (WI_{CDM-L}) values. L , a and b are used to designate measured values of three attributes of surface-color appearance as follows: L represents lightness, increasing from zero for black to 100 for perfect white; a represents redness when positive, greenness when negative, and zero for gray; and b represents yellowness
30 when positive, blueness when negative, and zero for gray. The concept of opponent colors was proposed by Hering in 1878. Starting in the 1940s, a number of measurable L , a , b dimensions have been defined by equations relating them to the basic CIE XYZ

tristimulus quantities defined in CIE Document No. 15. Measured values for a given color will depend on color space in which they are expressed [(TAPPI T 1213 sp-98 "Optical measurements terminology (related to appearance evaluation of paper")].

In one embodiment, the crosslinked fibers prepared with a polyol have a Whiteness Index ($WI_{(CDM-L)}$) value greater than about 69.0 when prepared under pilot plant conditions representative of commercial production. In another embodiment the crosslinked fibers have an L value greater than about 94.5. According to another embodiment the crosslinked fibers have an a value greater than about -1.55 and less than about -0.60. In yet another embodiment the crosslinked fibers have a b value less than about 8.50. The color properties of representative crosslinked cellulosic fibers are provided in Tables 1, 2. These fibers represent small scale tests (20 g. cellulose). Table 3 represents crosslinked fibers made in a pilot plant, representative of commercial production, using sorbitol as the polyol. Similar differences in color properties and brightness as those in Table 1 and 2 would be expected with C_4 - C_{12} polyols when processed under the pilot plant conditions. Whiteness Index is determined using a color difference meter (CDM) and is defined as:

$$WI_{(CDM-L)} = L - 3b.$$

Basic color measurement is made using commercially available instruments (e.g., Technibrite MicroTB-1C, Technydine Corp.). The instrument scans through the brightness and color filters. Fifty readings are taken at each filter position and averaged and the resulting values are printed out as Brightness, $R(X)$, $R(Y)$, and $R(Z)$. Brightness is ISO brightness (457 nm), $R(X)$ is absolute red reflectance (595 nm), $R(Y)$ is absolute green reflectance (557 nm), and $R(Z)$ is absolute blue reflectance (455 nm). The CIE tristimulus functions X , Y , and Z are then computed in accordance with the following equations: $X = 0.782 R(X) + 0.198 R(Z)$; $Y = R(Y)$; and $Z = 1.181 R(Z)$. Next L , a and b values are computed using the established equations (Technibrite Micro TB-1C Instruction Manual TTM 575-08, Oct. 30, 1989). $WI_{(CDM-L)}$ was calculated according to the equation: $WI_{(CDM-L)} = L - 3b$, according to TAPPI T 1216 sp-98 (TAPPI T 1216 sp-98 "Indices for whiteness, yellowness, brightness and luminous reflectance factor").

To further illustrate the principles, a discussion of whiteness and brightness is useful. Webster's Dictionary defines white as "the object color of greatest lightness characteristically perceived to belong to objects that reflect diffusely nearly all incident energy throughout the visible spectrum". Used as a noun or adjective, white is defined as

"free from color". Most natural and many man-made products are never "free from color". Whether the "white" product is fluff pulp, paper, textiles, plastics, or teeth, there is usually an intrinsic color, other than white, associated with it. Consider two hypothetical objects, the first that meets Webster's definition of white: one characterized by a flat spectrum of high reflectance and a second, which is the first with a small amount of blue colorant added (results in an unequal spectrum). Most people will judge the second as being the whiter of the two even though its total reflectance is lower in certain spectral regions. The first will be judged as a "yellow-white" while the second a "blue-white". Human color vision is more than just a sensation. It is also quite subjective and certain associations are unconsciously made. Blue-white is associated with "clean and pure", while "yellow-white" denotes "dirty, old or impure". The type and amounts of fillers and colorants to use, which hues are appropriate (e.g., red-blue, green-blue), and the optimal optical prescription to target have been the subject of considerable interest. The crosslinked cellulosic fibers prepared as described in Example 1, method A, without a polyol had brightness values significantly lower than the brightness of those crosslinked fibers where the cellulose fibers were treated with a crosslinking agent in the presence of a polyol. In contrast to the crosslinked cellulosic fibers prepared in the presence of the crosslinking agent and a polyol, all having a brightness greater than about 82.5 % ISO and $WI_{(CDM-L)}$ greater than about 73.77 by this method, the brightness achieved for the fibers crosslinked with only a crosslinking agent was as low as 79.2 % ISO and a $WI_{(CDM-L)}$ of 70.28.

In another aspect, the present application provides a method for making cellulosic fibers crosslinked in the presence of a polyol. In the method, cellulosic fibers are treated with an effective amount of a polyol in the presence of an effective amount of a crosslinking agent to achieve the brightness and color enhancements described herein. As used herein, an effective amount of crosslinking agent is from about 1 % to about 10 % by weight of the crosslinking agent based on the total weight of the cellulose fibers; an effective amount of the polyol is and from about 1 % to about 10 % by weight polyol based on the total weight of the fibers. In one embodiment, the fibers are treated with from about 1 % to about 10 % by weight of the polyol based on the weight of fibers. In another embodiment the fibers are treated with 2% to about 6% of the weight of the cellulose fiber with the polyol.

In yet another embodiment of the invention the fibers prepared with the crosslinking agent in the presence of a polyol have a wet bulk of at least about 16 cm³/g.

5 In another method the cellulose mat is treated with the polyol by methods known in the art, including spraying, rolling or dipping before the polyol treated sheet is impregnated with the crosslinking solution.

In another method the defiberized fiber is treated with the crosslinking agent is dried and the polyol is applied to the crosslinked treated fibers before the curing stage.

10 In general, the cellulose fibers may be prepared by a system and apparatus as described in U.S. Patent No. 5,447,977 to Young, Sr. et al. Briefly, the fibers are prepared by a system and apparatus that includes a conveying device for transporting a mat or web of cellulose fibers through a fiber treatment zone; an applicator for applying a treatment substance such as an aqueous solution of the crosslinking agent from a source to the fibers at the fiber treatment zone; a fiberizer for separating the individual cellulose fibers comprising the mat to form a fiber output comprised of substantially unbroken and
15 essentially singulated cellulose fibers; a dryer coupled to the fiberizer for flash evaporating residual moisture; and a controlled temperature zone for additional heating of fibers for drying and an oven for curing the crosslinking agent, to form dried and cured individualized crosslinked fibers.

20 As used herein, the term "mat" refers to any nonwoven sheet structure comprising cellulose fibers or other fibers that are not covalently bound together. The fibers include fibers obtained from wood pulp or other sources including cotton rag, hemp, grasses, cane, husks, cornstalks, or other suitable sources of cellulose fibers that may be laid into a sheet. The mat of cellulose fibers is preferably in an extended sheet form, and may be one of a number of baled sheets of discrete size or may be a continuous roll.

25 Each mat of cellulose fibers is transported by a conveying device, for example, a conveyor belt or a series of driven rollers. The conveying device carries the mats through the fiber treatment zone.

30 At the fiber treatment zone, an aqueous solution of the crosslinking agent is applied to the cellulose fibers. The crosslinking solutions are preferably applied to one or both surfaces of the mat using any one of a variety of methods known in the art, including spraying, rolling, or dipping. The polyol may be applied to the cellulose sheet before the application of the crosslinking solution, with the crosslinking solution, or after the

passage of the sheet through the fiberizer so that the polyol is applied to the individualized crosslinked treated fibers. In the latter case, the polyol can be injected into the hot air stream conveying the individualized fiber into the curing stage. Once the crosslinking solution and polyol have been applied to the mat, they may be uniformly distributed through the mat, for example, by passing the mat through a pair of rollers. After the fibers have been treated with the crosslinking agent and the polyol, the impregnated mat is fiberized by feeding the mat through a hammermill. The hammermill serves to disintegrate the mat into its component individual cellulose fibers, which are then air conveyed through a drying unit to remove the residual moisture. In one embodiment, the fibrous mat is wet fiberized.

The pulp is then air conveyed through an additional heating zone to bring the temperature of the pulp to the cure temperature. The cure temperature for citric acid is about 170°C. In one embodiment, the dryer comprises a first drying zone for receiving the fibers and for removing residual moisture from the fibers via a flash-drying method and a second heating zone for curing the crosslinking agent. Alternatively, in another embodiment, the treated fibers are blown through a flash-dryer to remove residual moisture, heated to a curing temperature, and then transferred to an oven where the treated fibers are subsequently cured. Overall, the treated fibers are dried and then cured for a sufficient time and at a sufficient temperature to effect crosslinking. Typically, the fibers are oven-dried and cured for about 15 seconds to about 20 minutes at a temperature from about 120°C to about 215°C.

As noted above, the present application relates to crosslinked cellulose fibers having improved brightness. Although available from other sources, cellulosic fibers useful for making crosslinked cellulosic fibers with improved color properties are derived primarily from wood pulp. Suitable wood pulp fibers can be obtained from well-known chemical processes such as the kraft and sulfite processes, with or without subsequent bleaching. The pulp fibers may also be processed by thermomechanical, chemithermomechanical methods, or combinations thereof. The pulp fiber is produced by chemical methods. Ground wood fibers, recycled or secondary wood pulp fibers, and bleached and unbleached wood pulp fibers can be used. The starting material is prepared from long-fiber coniferous wood species, such as southern pine, Douglas fir, spruce, and hemlock. Details of the production of wood pulp fibers are well-known to those skilled in the art.

These fibers are commercially available from a number of companies, including Weyerhaeuser Company. For example, suitable cellulose fibers produced from southern pine are available from Weyerhaeuser Company under the designations CF416, CF405, NF405, PL416, FR416, FR516, NB416, dissolving pulps from northern softwood include
5 MAC11 Sulfite, M919, WEYCELL and TR978 all of which have an alpha cellulose content of 95% and PH which has an alpha cellulose content of 91%. High purity mercerized pulps such as HPZ, HPZIII, HPZ4, and HPZ-XS available from Buckeye and Porosonier-J available from Rayonier are also suitable.

The wood pulp fibers can also be pretreated prior to use. This pretreatment may include
10 physical treatment, such as subjecting the fibers to steam or chemical treatment.

Although not to be construed as a limitation, examples of pretreating fibers include the application of fire retardants to the fibers, and surfactants or other liquids, such as solvents, which modify the surface chemistry of the fibers. Other pretreatments include incorporation of antimicrobials, pigments, and densification or softening agents. Fibers
15 pretreated with other chemicals, such as thermoplastic and thermosetting resins also may be used. Combinations of pretreatments also may be employed.

Method for determining fiber brightness. The brightness (% ISO) of cellulosic fibers crosslinked with citric acid was determined by TAPPI T 525 om-02.

The $WI_{(CDM-L)}$, brightness, L , a , and b values of representative crosslinked fibers prepared
20 with citric acid as the crosslinking agent, in the presence of various polyols and various levels of the polyol, using method A, are presented in Table 1; Table 2 represents fibers crosslinked with malic acid in the presence of sorbitol by the same method. Table 3 shows the effect of cure temperature and time on $WI_{(CDM-L)}$ and brightness when fibers are crosslinked with citric acid in the presence of sorbitol using the large scale production
25 method B.

The following examples are for the purposes of illustrating, and should not be construed as limitations.

Example 1

Representative Crosslinked Cellulosic Fibers Prepared With Sorbitol

30 In this example, methods for forming representative crosslinked fibers with improved brightness and color are described.

- Method A. A selected amount of a solution sufficient to apply 2, 8, and 2% by weight on cellulosic fibers, of sorbitol, citric acid and sodium hypophosphite, respectively, was applied to both sides of a twenty gram pulp sheet (CF416 or NF405, dried wood pulp fibers available from Weyerhaeuser Co.) using a 5 mL disposable syringe and 23.1 gauge
5 needle. The sample was held in a resealable plastic bag for 16-18 hours at room temperature, then broken into pieces (e.g., about 2x2 cm), passed through a laboratory fiberizer, and collected as a loose pad. The pad was broken into small pieces (e.g., about 3x3 cm), placed into a screen basket and cured at a fixed temperature and time in a Despatch V Series oven.
- 10 Citric acid crosslinked fibers with improved color and brightness properties prepared by this method with sorbitol and other polyols at 2 to 10 % of the weight of the cellulose fiber have $WI_{(CDM-L)}$ values, brightness, L , a , and b values described in Table 1; Table 2 represents brightness and color properties of fibers crosslinked with malic acid in the presence of sorbitol using the same method.
- 15 Method B. This pilot plant method is representative of commercial production. Pulp sheets in roll form (CF416, dried wood pulp fibers available from Weyerhaeuser Co.) were treated with citric acid and sorbitol according to the following procedure. The pulp sheet was fed from a roll through a constantly replenished bath of the crosslinking agent and sorbitol solution (i.e., an aqueous solution containing a citric acid and sorbitol
20 concentration determined by the weight add-on desired), then through a roll nip set to remove sufficient solution so that the pulp sheet after treating was at about 40 % by weight moisture content. The concentration of the bath was adjusted to achieve the desired level of chemical addition to the pulp sheet. After the roll nip, the wet sheet was fed through a hammermill to fiberize the pulp. The individualized fibers were then blown
25 through a flash dryer to affect drying and then to a cyclone where the treated cellulose fluff was separated from the air stream. The pulp was air conveyed through an additional heating zone to bring the temperature of the pulp to the cure temperature and then transferred to an oven where the treated fibers were subsequently cured. Crosslinked

fibers prepared by this method have the $WI_{(CDM-L)}$ brightness, L , a , and b values described in Table 3.

Method for determining fiber wet bulk. The wet bulk of cellulosic fibers crosslinked with citric acid and those crosslinked with citric acid in the presence of a polyol was determined by the Fiber Absorption Quality (FAQ) Analyzer (Weyerhaeuser Co. Federal Way, WA) using the following procedure.

In the procedure, a 4-gram sample of the pulp is put through a pinmill to open the pulp and then airlaid into a tube. The tube is then placed in the FAQ Analyzer. A plunger then descends on the fluff pad at a pressure of 0.6 kPa and the pad height bulk determined.

The weight is increased to achieve a pressure of 2.5kPa and the bulk recalculated. The result, the two bulk measurements on the dry fluff pulp at two different pressures. While under the 2.5 kPa pressure, water is introduced into the bottom of the tube (bottom of the pad). The time required for water to reach the plunger is measured. From this the absorption time and rate are determined. The final bulk of the wet pad at 2.5 kPa is also measured. The plunger is then withdrawn from the tube and the wet pad allowed to expand for 60 seconds. The plunger is reapplied at 0.6 kPa and the bulk determined. The final bulk of the wet pad at 0.6 kPa is considered the wet bulk (cm^3/g) of the pulp product. Fibers prepared by method B using citric acid and sorbitol have a bulk of at least 16.1.

Table 1

Properties of Representative Crosslinked Fibers Prepared By Treating Cellulose Fibers With Citric Acid In The Presence Of Various Polyols

Pulp	Additive	Additive Wt. % on fiber	FAQ Wet Bulk cc/g	ISO Brightness %	L	a	b	$WI_{(CDM-L)}$
NF405	No additive	0	18.5	82.5	96.13	-1.53	7.68	73.09
NF405	Erythritol	2	18.6	84.6	95.88	-1.32	5.69	78.81
NF405	Xylitol	2	18.3	85.2	96.04	-1.27	5.47	79.63
NF405	Arabinitol	2	19	85.4	96	-1.09	5.32	80.04
NF405	Ribitol	2	18.7	85.7	96.15	-1.05	5.33	80.16
NF405	Sorbitol (Glucitol)	2	18.5	85.9	96.2	-1.26	5.12	80.84
NF405	Mannitol	2	19.4	85.2	95.96	-1.29	5.38	79.82
NF405	Lactitol	2	19	84	96.06	-1.29	6.44	76.77
NF405	Maltitol	2	19.1	84.2	96.05	-1.3	6.29	77.18

Pulp	Additive	Additive Wt. % on fiber	FAQ Wet Bulk cc/g	ISO Brightness %	L	a	b	WI _(CDM-L)
NF405	Isomalt	2	18.7	84	95.96	-1.09	6.29	77.09
NF405	myo-Inositol	2	19	84.1	95.94	-1.22	6.19	77.37
CF416	No additive	0	17.6	82.1	95.58	-1.38	7.27	73.77
CF416	Erythritol	2	17.9	84.5	95.35	-1.12	5.01	80.32
CF416	Xylitol	2	17.7	84.8	95.45	-1.11	4.97	80.54
CF416	Arabinitol	2	18.2	84.7	95.48	-0.92	5.08	80.24
CF416	Ribitol	2	18.4	85.1	95.53	-0.91	4.88	80.89
CF416	Sorbitol (Glucitol)	2	17.6	85	95.44	-1.07	4.78	81.1
CF416	Mannitol	2	17.9	85.3	95.5	-0.95	4.59	81.73
CF416	Lactitol	2	18.4	83.6	95.47	-1.05	5.92	77.71
CF416	Maltitol	2	17.9	84.5	96.14	-1.27	6.14	77.72
CF416	Isomalt	2	17.7	81.7	94.71	-0.84	6.3	75.81
CF416	myo-Inositol	2	18.6	83.4	95.49	-1.09	6.11	77.16
NF405	No additive	0	18.5	82.5	96.13	-1.53	7.68	73.09
NF405	Erythritol	4	17.8	86.7	96.3	-1.03	4.7	82.2
NF405	Xylitol	4	17.4	86.5	96.15	-1.15	4.67	82.14
NF405	Arabinitol	4	18.3	86.5	96.24	-1	4.82	81.78
NF405	Ribitol	4	18.7	86.2	96.09	-1.05	4.82	81.63
NF405	Sorbitol (Glucitol)	4	17.5	86.1	96.08	-1.18	4.86	81.5
NF405	Mannitol	4	18.3	86.2	96.18	-1.11	4.89	81.51
NF405	Lactitol	4	18.6	85.1	96.3	-1.14	5.9	78.6
NF405	Maltitol	4	18.5	85.4	96.01	-1.1	5.32	80.05
NF405	Isomalt	4	18.3	84.9	96.2	-1.1	5.95	78.35
NF405	myo-Inositol	4	18.5	84.7	95.82	-1.12	5.54	79.2
CF416	No additive	0	17.6	82.1	95.58	-1.38	7.27	73.77
CF416	Erythritol	4	16.8	85.4	95.3	-0.94	4.26	82.52
CF416	Xylitol	4	16.5	85.8	95.58	-0.9	4.32	82.62
CF416	Arabinitol	4	17.4	86.7	95.89	-0.69	4.17	83.38
CF416	Ribitol	4	17.5	86.1	95.63	-0.72	4.21	83
CF416	Sorbitol (Glucitol)	4	17.2	85.7	95.43	-0.96	4.2	82.83
CF416	Mannitol	4	17.8	85.3	95.43	-1	4.55	81.78
CF416	Lactitol	4	17.8	84.5	95.7	-0.98	5.59	78.93
CF416	Maltitol	4	17	85.4	95.76	-0.8	4.9	81.06
CF416	Isomalt	4	17.2	82.7	94.8	-0.79	5.63	77.91
CF416	myo-Inositol	4	17.7	84.58	95.5	-0.9	5.3	79.6
NF405	No additive	0	18.5	82.5	96.13	-1.53	7.68	73.09
NF405	Erythritol	6	16.9	86.4	95.92	-1.09	4.4	82.72
NF405	Xylitol	6	17.0	87	96.19	-1.12	4.35	83.14
NF405	Arabinitol	6	17.1	86.9	96.25	-0.96	4.52	82.69
NF405	Ribitol	6	17.9	86.9	96.32	-0.92	4.65	82.37
NF405	Sorbitol (Glucitol)	6	17.3	86.6	96.09	-1.12	4.47	82.68
NF405	Mannitol	6	17.7	86.3	95.99	-1.18	4.58	82.25
NF405	Lactitol	6	18.3	85.5	96.43	-1.11	5.79	79.06
NF405	Maltitol	6	17.6	85.7	96.41	-1.09	5.58	79.67
NF405	Isomalt	6	17.7	85.1	96.03	-1.06	5.53	79.44
NF405	myo-Inositol	6	17.9	85.2	96.08	-1.05	5.52	79.52
CF416	No additive	0	17.6	82.1	95.58	-1.38	7.27	73.77

Pulp	Additive	Additive Wt. % on fiber	FAQ Wet Bulk cc/g	ISO Brightness %	L	a	b	WI _(CDM-L)
CF416	Erythritol	6	16.2	86.4	95.8	-0.92	4.24	83.08
CF416	Xylitol	6	16	86.4	95.67	-0.86	3.99	83.7
CF416	Arabinitol	6	16.4	86.6	95.61	-0.62	3.83	84.12
CF416	Ribitol	6	17.2	86.4	95.6	-0.69	3.91	83.87
CF416	Sorbitol (Glucitol)	6	16.1	85.9	95.42	-0.83	4.05	83.27
CF416	Mannitol	6	16.9	86.3	95.62	-0.78	4.02	83.56
CF416	Lactitol	6	17.4	84.8	95.59	-0.83	5.12	80.23
CF416	Maltitol	6	16.9	82.2	94.41	-0.69	5.43	78.12
CF416	Isomalt	6	16.5	82.8	94.92	-0.75	5.65	77.97
CF416	myo-Inositol	6	16.8	84.4	95.3	-0.71	5.03	80.21
NF405	No additive	0	18.5	82.5	96.13	-1.53	7.68	73.09
NF405	Erythritol	8	16.1	86.5	95.67	-0.82	3.96	83.79
NF405	Xylitol	8	16.5	87	96.25	-1.13	4.42	82.99
NF405	Arabinitol	8	-	-	-	-	-	-
NF405	Ribitol	8	-	-	-	-	-	-
NF405	Sorbitol (Glucitol)	8	16.1	86.9	96.23	-1.13	4.42	82.97
NF405	Mannitol	8	16.8	87.2	96.14	-0.93	4.09	83.87
NF405	myo-Inositol	8	-	-	-	-	-	-
CF416	No additive	0	17.6	82.1	95.58	-1.38	7.27	73.77
CF416	Erythritol	8	15.2	86.9	96.29	-1.07	4.54	82.67
CF416	Xylitol	8	15.6	86.8	95.8	-0.85	3.88	84.16
CF416	Arabinitol	8	-	-	-	-	-	-
CF416	Ribitol	8	-	-	-	-	-	-
CF416	Sorbitol (Glucitol)	8	15.7	86.7	95.68	-0.86	3.81	84.25
CF416	Mannitol	8	16.2	86.6	95.52	-0.74	3.63	84.63
CF416	myo-Inositol	8	-	-	-	-	-	-
NF405	No additive	0	18.5	82.5	96.13	-1.53	7.68	73.09
NF405	Erythritol	10	15.6	86.8	95.58	-0.61	3.58	84.84
NF405	Xylitol	10	15.8	87.1	96.27	-1.13	4.34	83.25
NF405	Arabinitol	10	-	-	-	-	-	-
NF405	Ribitol	10	-	-	-	-	-	-
NF405	Sorbitol (Glucitol)	10	15.9	87.2	96.19	-1.08	4.17	83.68
NF405	Mannitol	10	16.2	87.3	96.22	-1.09	4.17	83.71
NF405	myo-Inositol	10	-	-	-	-	-	-
CF416	No additive	0	17.6	82.1	95.58	-1.38	7.27	73.77
CF416	Erythritol	10	14.9	86.6	95.63	-1	3.77	84.32
CF416	Xylitol	10	14.7	87.1	95.75	-0.81	3.56	85.07
CF416	Arabinitol	10	-	-	-	-	-	-
CF416	Ribitol	10	-	-	-	-	-	-
CF416	Sorbitol (Glucitol)	10	15	87	95.77	-0.79	3.66	84.79
CF416	Mannitol	10	15.7	86.2	95.53	-0.84	3.94	83.71
CF416	myo-Inositol	10	-	-	-	-	-	-

Experimental conditions: 8% by weight citric acid on cellulose fibers, 2 % by weight sodium hypophosphite on cellulose fibers, additive as listed, cured at 170°C for 7 min.

Table 2

Properties Of Representative Crosslinked Fibers Prepared By Treating Cellulose Fibers
With Malic Acid In The Presence Of Sorbitol

Pulp	Crosslinking Agent	Additive	Additive, Wt. % on fiber	FAQ Wet Bulk, cc/g	ISO Brightness, %	L	a	b	WI _(CDM-L)
NF 405	Malic Acid	None	-	17.3	81.1	95.53	-1.2	7.88	71.89
NF405	Malic Acid	Sorbitol	2	16.4	83.9	96.04	-1.14	6.48	76.6
NF405	Malic Acid	Sorbitol	4	15.8	84.7	96.24	-1.14	6.17	77.73
NF405	Malic Acid	Sorbitol	6	14.7	85.5	96.37	-1.08	5.76	79.09
CF416	Malic Acid	None	-	16.8	79.2	94.64	-1.12	8.12	70.28
CF416	Malic Acid	Sorbitol	2	15.7	84.2	95.71	-0.95	5.81	74.28
CF416	Malic Acid	Sorbitol	4	14.9	84	95.47	-1	5.66	78.49
CF416	Malic Acid	Sorbitol	6	13.8	85.9	96.03	-0.83	4.92	81.27

5 Experimental conditions: 8% by weight malic acid on cellulose fibers, 2 % by weight sodium hypophosphite on cellulose fibers, sorbitol as listed, cured at 170°C for 7 min.

Table 3

Properties Of Representative Crosslinked Fibers Prepared By Treating Cellulose Fibers
With Citric Acid In The Presence Of Sorbitol And Cured Under Different Conditions

10

Pulp	Additive	Additive, Wt. % on fiber	Cure Temp, °F	Cure Time Min.	FAQ Wet Bulk, cc/g	ISO Brightness, %	L	a	b	WI _(CDM-L)
CF416	none	0	360	5	16.2	78.4	94.7	-1.58	8.77	68.39
CF416	Sorbitol	1.5	360	5	16.1	83.09	95.25	-1.28	6.0	77.25
CF416	none	0	360	7	17.2	75.6	94.1	-1.59	10.11	63.77
CF416	Sorbitol	1.5	360	7	16.6	82.7	95.8	-1.54	7.02	74.74
CF416	none	0	380	5	17.5	74.5	93.9	-1.57	10.67	61.89
CF416	Sorbitol	1.5	380	5	16.6	81.69	95.36	-1.31	7.21	73.73
CF416	none	0	380	7	17.7	70.3	92.8	-1.49	12.48	55.36
CF416	Sorbitol	1.5	380	7	16.8	79.50	94.96	-1.51	8.33	69.97

Experimental conditions: 6% by weight citric acid on cellulose fibers, 0.75 % by weight sodium hypophosphite on cellulose fibers, additive as listed, cured as indicated

The present application provides cellulosic fibers having improved brightness and color.

15 The fibers are intrafiber crosslinked cellulosic fibers obtainable from cellulosic fibers by treatment with a crosslinking agent in the presence of a polyol. The crosslinked fibers can be formed from cellulosic fibers by treatment with an amount of polyol in the presence of a crosslinking agent effective to provide the color and brightness enhancement described herein.

The crosslinked cellulosic fibers with improved color properties can be incorporated into an absorbent product. Such products can further include other fibers such as fluff pulp fibers, synthetic fibers, other crosslinked fibers, and absorbent materials such as superabsorbent polymeric materials. Representative absorbent products that can include the fibers include infant diapers, adult incontinence products, and feminine hygiene products. The fibers can be included in liquid acquisition, distribution, or storage layers. The crosslinked cellulosic fibers with improved color properties can also be incorporated into tissue and towel products.

Additionally, the fibers can be incorporated into paperboard products, including single and multi-ply paperboard products. Paperboard products that include the fibers can be used in insulation applications, for example, insulated cups and containers. Paperboard products that include the fibers can also be used as packaging materials.

While various embodiments of the invention have been illustrated and described, it will be appreciated that the present invention can be practiced by other than the described embodiments, which are presented for purposes of illustration and not limitation, and present invention is limited only by the claims that follow.